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Substitute for form 1449/PTO		Complete If Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)		Application Number	10/552,314-Conf. #1865
		Filing Date	September 5, 2006
		First Named Inventor	Annie BARDAT
		Art Unit	1644
		Examiner Name	Y. Kim
Sheet	1	of	2
		Attorney Docket Number	0040-0158PUS1

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Document Number Number/Kind Code ² (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
	AA*	US-5,945,098	08-31-1999	Sarno et al.	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Foreign Patent Document Country Code ³ -Number/Kind Code ² (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁴
	BA	WO-97/04801-A1	02-13-1997	GENENTECH INC		
	BB	WO-02/092632	11-21-2002	LAB FRANCAIS DU FRACTIONNEMENT		
	BC	WO-2002/013860	02-21-2002	CHUGAI PHARMACEUTICAL CO LTD		✓
	BD	JP-61-191622	08-26-1986	Green Cross Corp		✓
	BE	EP-0196761-A2	10-08-1986	GREEN CROSS CORP		
	BG	JP-5-025058	02-02-1993	HAGIWARA YOSHIHIDE		✓
	BH	EP-0597101	05-18-1994	HAGIWARA HIDEAKI		
	BI	JP-63-088197	04-19-1988	Tosoh Corp		ABS
	BJ	JP-9-500894	01-28-1997			ABS
	BK	WO-98/44948-A2	10-15-1998	CANGENE CORP		
	BL	JP-11-510170	09-07-1999			✓
	BL	EP-1314437	05-28-2003	CHUGAI PHARMACEUTICAL CO LTD		

Examiner Signature	Date Considered
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NON PATENT LITERATURE DOCUMENTS			
Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
	CA	BLEEKER, Wim K. et al., "Vasoactive side effects of intravenous immunoglobulin preparations in a rat model and their treatment with recombinant platelet-activating factor acetylhydrolase," <i>Blood</i> , March 1, 2000, Vol. 95, No. 5, pp. 1856-1861.	
	CB	PIKAL, Michael J., "Freeze-drying of proteins, part II: Formulation selection," <i>BioPharm</i> , October 1990, Vol. 3, No. 9, pp. 26-30.	
	CC	ARAKAWA, Tsutomu et al., "Protein-solvent interactions in pharmaceutical formulations," <i>Pharmaceutical Research</i> , 1991, Vol. 8, No. 3, pp. 285-291.	
	CD	OSTERBERG, Thomas et al., "Development of a freeze-dried albumin-free formulation of recombinant factor VIII SQ," <i>Pharmaceutical Research</i> , 1997, Vol. 14, No. 7, pp. 892-898.	
	CE	GUO, Wei et al., "Raman evidence that the lyoprotectant poly(ethylene glycol) does not restore nativity to the heme active site of horseradish peroxidase suspended in organic solvents," <i>Biomacromolecules</i> , 2002, Vol. 3, No. 4, pp. 846-849.	
	CF	CHIDWICK, K. et al., "Clinical experience with a new solvent detergent-treated intravenous immunoglobulin free of hypotensive effects," <i>Vox Sanguinis</i> , 1999, Vol. 77, No. 4, pp. 204-209.	
	CG	COHN, E. J. et al., "Preparation and properties of serum and plasma proteins. IV. A system for the separation into fractions of the protein and lipoprotein components of biological tissues and fluids," <i>J. Am. Chem. Soc.</i> , March 1946, Vol. 68, pp. 459-475.	
	CH	KISTLER, P. et al., "Large scale production of human plasma fractions. Eight years experience with the alcohol fractionation procedure of Nitschmann, Kistler and Lergier," <i>Vox Sang.</i> , 1962, Vol. 7, pp. 414-424.	
	CI	STEINBUCH, M. et al., "Isolément de l'immunoglobuline IgG du plasma humain a l'aide de l'acide caprylique," <i>Rev. Fr. Etud. Clin. Biol.</i> , Dec. 1969, Vol. 14, No. 10, pp. 1054-1058.	
	CJ	FERNANDES, Peter M. et al., "Preparation of a stable intravenous gamma-globulin: Process design and scale-up," <i>Vox Sang.</i> , 1980, Vol. 39, No. 2, pp. 101-112.	
	CK	LEVINE, Howard L. et al., "The use of surface tension measurements in the design of antibody-based product formulations," <i>Journal of Parenteral Science & Technology</i> , 1991, Vol. 45, No. 3, pp. 160-165.	
	CL	Pharmacopée Européenne, 4ème édition, chap. <<Immunoglobuline humaine normale pour administration par voie intraveineuse>>, Methode 2.6.17.	

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